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CRF Design Helper

Healthcare

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This guide explains what the tool produces, how to use the output, and what to watch for. The actual output (draft/report) is generated on the tool page.

Background

Helps design Case Report Forms — modules, fields and edit checks aligned to the protocol and CDASH principles. Use this for CRF design; for the protocol synopsis use the Study Protocol Synopsis.

Purpose of this tool & output

Design case report forms for a study

How the output is used

Use the helper to design case report forms that capture the study data cleanly and per protocol.

Important cautions

AI-generated design aid; a data-management/clinical expert must finalise.

Companion documents

- Clinical Trial Protocol Review
- Informed Consent Form Reviewer
- GCP Compliance Checklist
- Ethics Committee Submission Guide

This is AI-assisted guidance. Verify with a qualified professional before any binding decision. Fees, authorities and processes vary by state and change over time — confirm with your state authority.